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AMENDMENTS TO THE CLAIMS

1. (Currently Amended) A hypodermic needle assembly for use in making

intradermal injections, comprising:

a hub portion that is able to be attached to a drug container;

a needle supported by the hub portion, the needle having a hollow body with a

forward end extending away from the hub portion; and

a limiter portion that is proximate to the needle and surrounds the needle and

extends away from the hub portion toward the forward end of the

needle, the limiter portion is non movable with respect to said hub

portion and said limiter having a skin engaging surface that is adapted

to be received against skin of an animal to receive an intradermal

injection, the needle forward end extending beyond the skin engaging

surface a preselected distance which is set during manufacture of the

needle assembly such that the limiter portion limits an amount that the

needle is able to penetrate through the skin of an animal.

2. (Original) The assembly of claim 1, wherein the hub portion and the limiter

portion are integrally formed as a single piece made from a plastic material.

3. (Original) The assembly of claim 1, wherein the hub portion and the limiter

portion are formed as separate pieces.

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4. (Original) The assembly of claim 3, wherein the limiter portion includes an inner

cavity that receives at least a portion of the hub portion and the inner cavity includes an

abutment surface that engages corresponding structure on the hub portion to thereby limit

the amount that the needle forward end extends beyond the skin engaging surface.

5. (Original) The assembly of claim 3, wherein the limiter portion is integrally

formed as part of the syringe and the hub portion is received within the limiter portion.

6. (Original) The assembly of claim 5, wherein the skin engaging surface

surrounds the needle, and has a thickness defined between an inner diameter and an

outer diameter and wherein the inner diameter is at least five times greater than an outside

diameter of the needle.

7. (Original) The assembly of claim 6, wherein the skin engaging surface is

generally circular.

8. (Original) The assembly of claim 1, wherein the skin engaging surface includes

a central opening that is slightly larger than an outside dimension of the needle and the

skin engaging surface is continuous.

9. (Original) The assembly of claim 1, wherein the skin engaging surface is

generally flat and extends through a plane that is generally perpendicular to an axis of the

needle.

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10. (Original) The needle assembly of claim 1, wherein the selected distance that

the forward end of the needle extends beyond the skin engaging surface is fixed.

11. (Original) The assembly of claim 1, wherein the selected distance is in the

range from approximately .5mm to approximately 3mm.

12. (Original) The assembly of claim 1, wherein the skin engaging surface includes

a contact surface area that is large enough to stabilize the assembly in a desired

orientation relative to the skin.

13. (Original) The assembly of claim 12, wherein the desired orientation is generally

perpendicular to the skin.

14. (Original) The assembly of claim 1, wherein the drug container is a syringe and

the animal is human.

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15. (Currently Amended) An intradermal delivery device for use in making

intradermal injections, comprising:

a drug container having a reservoir adapted to contain a selected substance and

an outlet port that allows the substance to exit the reservoir during an

injection;

a needle in fluid communication with the outlet port, the needle having a forward

end that is adapted to penetrate an the skin of an animal; and

a limiter that surrounds the needle and is fixed with respect to said outlet port and

said limiter has a skin engaging surface that is adapted to be placed

against the skin of the animal to receive an intradermal injection, the

needle forward end extending away from the skin engaging surface a

preselected distance which is set during manufacture of the

intradermal delivery device such that the limiter limits an amount that

the needle forward end penetrates the skin.

16. (Original) The device of claim 15, wherein the drug container is a syringe

including a generally hollow, cylindrical body portion and a plunger that is received within

the reservoir, the plunger being selectively movable within the reservoir to cause the

substance to be forced out of the outlet port during an injection.

17. (Original) The device of claim 15, including a hub portion that supports the

needle and the hub portion is selectively secured to the drug container near the outlet port.

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Claims 18-24 Cancelled

25. (Original) The device of claim 15, wherein the needle has a length and wherein

the selected distance is much less than the needle length.

26. (Original) The device of claim 25, wherein the selected distance is fixed and is

in the range from approximately .5mm to approximately 3mm.

27. (Original) The device of claim 15, wherein the skin engaging surface is

generally flat and extends through a plane that is generally perpendicular to an axis of the

needle.

28. (Original) The device of claim 15, wherein the skin engaging surface includes a

central opening that is slightly larger than an outside dimension of the needle and the skin

engaging surface is continuous.

29. (Original) The device of claim 15, wherein the skin engaging surface includes a

contact surface area that is large enough to stabilize the assembly in a desired orientation

relative to the skin.

30. (Original) The device of claim 15, wherein the desired orientation is generally

perpendicular to the skin.

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31. (Original) The device of claim 15, wherein the drug container is prefilled with

a substance.

32. (Previously Presented) A method of intradermally injecting at least one

substance such as a drug, vaccine or the like into the skin, comprising the steps of:

pressing a needle perpendicularly to the skin of the animal to receive an injection,

said needle in fluid communication with an outlet port of a drug container

having a reservoir adapted to contain a selected substance and the outlet

port allows the substance to exit the reservoir during an intradermal

injection;

injecting the substance into the skin of the animal with the depth of penetration of

the needle being limited to the intradermal space by a limiter that surrounds

the needle and is fixed with respect to said outlet port and said limiter and

has a skin engaging surface that is adapted to be placed against the skin of

the animal and a forward end of the needle extending away from the skin

engaging surface a preselected distance which is set during manufacture

of the needle and limiter such that the limiter limits an amount that the

needle forward end penetrates the skin of the animal.

33. (Original) The method of claim 32, wherein the step of pressing the needle

perpendicularly to the skin of the animal includes orienting the needle perpendicularly to

the skin.

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34. (Original) The method of claim 32, wherein the step of injecting the substance

includes moving a plunger that is received within the reservoir, with the plunger being

selectively movable within the reservoir to cause the substance to be forced out of the

outlet port during the injection.

35. (Canceled)

36. (Original) The method of claim 32, further comprising the step of filling the drug

container with the substance to be intradermally injected.

37. (Original) The method of claim 32, wherein said drug container is a syringe and

said animal is human.

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38. (Currently Amended) An intradermal delivery device for use in making

intradermal injections, comprising:

a drug container formed of glass having a reservoir adapted to contain a selected

substance and an outlet port that allows the substance to exit the reservoir

during an injection;

a needle in fluid communication with the outlet port, the needle having a forward end

that is adapted to penetrate an the skin of an animal; and

an limiter integrally formed on said drug container that is substantially proximate to the

needle and said limiter surrounds the needle and is fixed with respect to said

outlet port and said limiter has a skin engaging surface that is adapted to be

placed against the skin of the animal to receive an intradermal injection, the

needle forward end extending away from the skin engaging surface a

preselected distance which is set during manufacture of the intradermal

delivery device such that the limiter limits an amount that the needle forward

end penetrates the skin.

39. (Previously Presented) The device of claim 38, wherein the drug container is a

syringe including a generally hollow, cylindrical body portion and a plunger that is received

within the reservoir, the plunger being selectively movable within the reservoir to cause the

substance to be forced out of the outlet port during an injection.

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40. (Previously Presented) The device of claim 38, including an integrally formed

hub portion that supports the needle and the hub portion is secured to the drug container

near the outlet port.

41. (Previously Presented) The device of claim 38, wherein the needle has a length

and wherein the preselected distance is much less than the needle length.

42. (Previously Presented) The device of claim 41, wherein the selected distance is

fixed and is in the range from approximately .5mm to approximately 3mm.

43. (Previously Presented) The device of claim 38, wherein the skin engaging

surface is generally flat and extends through a plane that is generally perpendicular to an

axis of the needle.

44. (Previously Presented) The device of claim 38, wherein the skin engaging

surface includes a central opening that is slightly larger than an outside dimension of the

needle and the skin engaging surface is continuous.

45. (Previously Presented) The device of claim 38, wherein the skin engaging

surface includes a contact surface area that is large enough to stabilize the assembly in a

desired orientation relative to the skin.

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46. (Previously Presented) The device of claim 38, wherein the desired orientation is generally perpendicular to the skin.

47. (Previously Presented) The device of claim 38, wherein the drug container is prefilled with a substance.